

$$\therefore \dot{V}_m = 110 \text{ cm}^3 \times \left( \frac{1 \text{ m}}{100 \text{ cm}} \right)^3 = 1.1 \times 10^{-4} \text{ m}^3$$

$$\rho_m = 2,700 \text{ kg/m}^3$$

$$\rho_{\text{env}} = 790 \text{ kg/m}^3$$

a)



$$\vec{F}_{\text{net},y} = m \vec{a}_y$$

$$\text{so } T + F_b - m_m g = 0$$

$$T = m_m g - F_b$$

$$\text{now } F_b = \rho_{\text{env}} \dot{V}_m g = \rho_{\text{env}} \dot{V}_m g = m_{\text{env}} g$$

now

$$m_m = \rho_m \dot{V}_m = \left( 2,700 \frac{\text{kg}}{\text{m}^3} \right) (1.1 \times 10^{-4} \text{ m}^3) = 0.297 \text{ kg}$$

$$m_{\text{env}} = \rho_{\text{env}} \dot{V}_m = \left( 790 \frac{\text{kg}}{\text{m}^3} \right) (1.1 \times 10^{-4} \text{ m}^3) = 0.0869 \text{ kg}$$

$$T = (m_m - m_{\text{env}})g = (0.297 \text{ kg} - 0.0869 \text{ kg})(9.8 \text{ m/s}^2)$$

$$[T = 2.1 \text{ N}]$$

2.  $n = 0.10 \text{ mol}$  of ARGON

$$V_i = 55 \text{ cm}^3 \times \left(\frac{1 \text{ m}}{100 \text{ cm}}\right)^3 = 5.5 \times 10^{-5} \text{ m}^3$$

$$\text{now } pV = nRT$$

$$T_i = 296 \text{ K}$$

$$V_f = 220 \text{ cm}^3 = 2.2 \times 10^{-4} \text{ m}^3$$

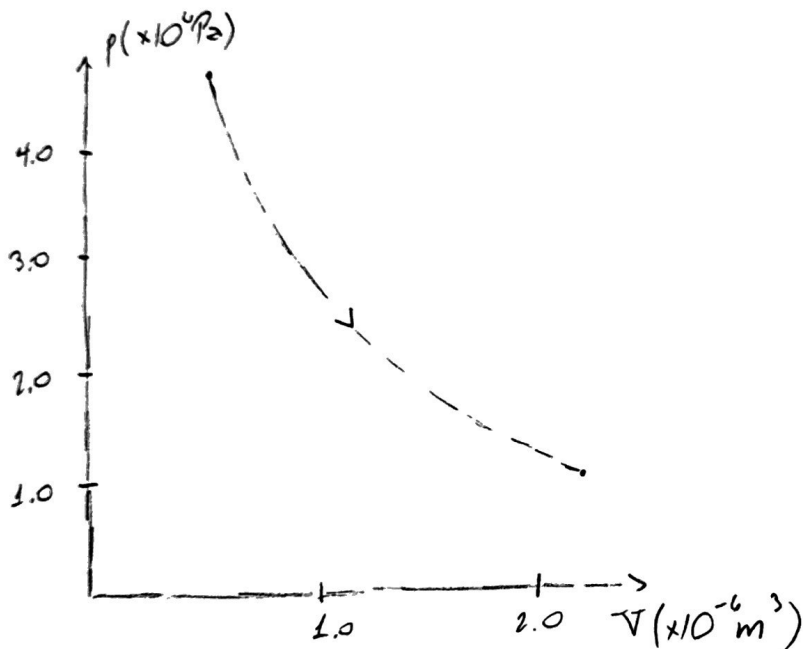
INITIAL

$$a) \quad p_i = \frac{nRT_i}{V_i} = \frac{(0.10 \text{ mol})(8.315 \text{ J/mol}\cdot\text{K})(296 \text{ K})}{(5.5 \times 10^{-5} \text{ m}^3)} = 4.47 \times 10^6 \text{ Pa}$$

ISOTHERMAL EXPANSION  $T_f = T_i$  so

$$p_i V_i = p_f V_f \Rightarrow p_f = p_i \frac{V_i}{V_f} = (4.47 \times 10^6 \text{ Pa}) \frac{(5.5 \times 10^{-5} \text{ m}^3)}{(2.2 \times 10^{-4} \text{ m}^3)}$$

$$[p_f = 1.11 \times 10^6 \text{ Pa}]$$



$$P_1 = 3.03 \times 10^5 \text{ Pa} = P_2$$

$$V_1 = 100 \text{ cm}^3 \times \left(\frac{1 \text{ m}}{100 \text{ cm}}\right)^3 = 1.0 \times 10^{-4} \text{ m}^3$$

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2} = \frac{P_3 V_3}{T_3}$$

$$V_2 = 300 \text{ cm}^3 = 3.0 \times 10^{-4} \text{ m}^3 = V_3 = 3V_1$$

$$T_1 = T_3 = 100^\circ \text{C} = 373 \text{ K}$$

Question: States 1 to 2, isobaric process so  $P_1 = P_2$

$$\frac{V_1}{T_1} = \frac{V_2}{T_2} \therefore T_2 = T_1 \frac{V_2}{V_1} = 373 \text{ K} \left(\frac{3V_1}{V_1}\right) = 1,120 \text{ K}$$

$$T_1 = T_3 = 373 \text{ K}$$

$$T_2 = 1,120 \text{ K}$$

For process 1 to 2, isobaric process...

$$\Delta E_{\text{th}} = n C_V \Delta T = \quad \text{now } PV = nRT \text{ so } n = \frac{PV}{RT} = \frac{(3.03 \times 10^5 \text{ Pa})(1.0 \times 10^{-4} \text{ m}^3)}{(8.31 \text{ J/mol} \cdot \text{K})(373 \text{ K})}$$

$$= (9.78 \times 10^{-3} \text{ mol}) \left(12.5 \frac{\text{J}}{\text{mol} \cdot \text{K}}\right) (1,120 \text{ K} - 373 \text{ K}) = 9.78 \times 10^{-3} \text{ mol}$$

$$[\Delta E_{\text{th}} = 91 \text{ J}]$$

$$W = -p \Delta V = -(3.03 \times 10^5 \text{ Pa})(3.0 \times 10^{-4} \text{ m}^3 - 1.0 \times 10^{-4} \text{ m}^3) = -61 \text{ J}$$

$$[W = -61 \text{ J}]$$

$$\text{now } \Delta E_{\text{th}} = Q + W \text{ so } Q = \Delta E_{\text{th}} - W = 91 \text{ J} - (-61 \text{ J}) = 152 \text{ J}$$

$$[Q = 150 \text{ J}]$$

For process 2 to 3, isochoric process  $\therefore W = 0 \text{ J}$

$$\Delta E_{\text{th}} = n C_V \Delta T = (9.78 \times 10^{-3} \text{ mol}) (12.5 \text{ J/mol} \cdot \text{K}) (373 \text{ K} - 1,120 \text{ K}) = -91 \text{ J}$$

$$\text{so } \Delta E_{\text{th}} = W + Q = Q$$

$$[\Delta E_{\text{th}} = Q = -91 \text{ J}] \therefore [W = 0 \text{ J}]$$

$$4. m_{ce} = 0.512 \text{ kg}$$

$$T_{i,ce} = 190^\circ\text{C} = 1,063 \text{ K}$$

$$m_{H_2O} = 0.210 \text{ kg}$$

$$T_{i,H_2O} = 22^\circ\text{C} = 295 \text{ K}$$

$$T_{f,ce} = T_{f,H_2O} = 100^\circ\text{C} = 373 \text{ K}$$

$$V_{H_2O} = 210 \times 10^{-3} \text{ L} \times \frac{1 \text{ m}^3}{1000 \text{ L}} = 2.1 \times 10^{-4} \text{ m}^3$$

$$m = \rho_{H_2O} V_{H_2O} = (1,000 \text{ kg/m}^3)(2.1 \times 10^{-4} \text{ m}^3) = 0.210 \text{ kg}$$

$$Q_{ce} + Q_{H_2O} = 0$$

$$m_{ce} C_{ce} (373 \text{ K} - 1063 \text{ K}) + m_{H_2O} C_{H_2O} (373 \text{ K} - 295 \text{ K}) + m_{H_2O}^* L_{v,H_2O} = 0$$

so

$$m_{H_2O}^* = \frac{-m_{ce} C_{ce} (-690 \text{ K}) - m_{H_2O} C_{H_2O} (78 \text{ K})}{L_{v,H_2O}}$$

$$= \frac{-(0.512 \text{ kg})(449 \text{ J/kg} \cdot \text{K})(-690 \text{ K}) - (0.210 \text{ kg})(4,190 \text{ J/kg} \cdot \text{K})(78 \text{ K})}{(22.6 \times 10^5 \text{ J/kg})}$$

$$= \frac{1.586 \times 10^5 \text{ J} - 6.86 \times 10^4 \text{ J}}{22.6 \times 10^5 \text{ J/kg}} = 3.98 \times 10^{-2} \text{ kg} = 39.8 \text{ g}$$

so the fraction of the  $H_2O$  that boils is...

$$\frac{m_{H_2O}^*}{m_{H_2O}} = \frac{39.8 \text{ g}}{210 \text{ g}} = 0.19$$